

## Claims

What is claimed is:

1. A surface plasmon resonance system, comprising:  
a support comprising a metal film, a first medium adjacent a first  
5 side of the metal film and a second medium adjacent a second side of the metal  
film;  
an illumination source operable to illuminate at least one incident  
region of the metal film through the first medium;  
at least one assay region on the second side of the metal film  
10 opposing the incident region; and  
a collector operable to collect illumination reflected from the metal  
film;  
wherein at least one of the illumination source and the collector  
comprise an optical integrated circuit.  
15
2. The surface plasmon resonance system according to claim 1,  
wherein the optical integrated circuit comprises at least one arrayed waveguide  
grating spectrometer.
- 20 3. The surface plasmon resonance system according to claim 1,  
wherein the second medium comprises a first member of a binding pair and a  
second member of a binding pair.
4. The surface plasmon resonance system according to claim 1,  
25 wherein the metal film comprises at least about 100 incident regions and at least  
about 100 corresponding assay regions, the illumination source operable to  
illuminate the at least about 100 incident regions.
5. The surface plasmon resonance system according to claim 1,

LWBA101USA

wherein the metal film is comprised within one of a gene chip, a DNA chip, or a protein chip.

5           6.     The surface plasmon resonance system according to claim 3,  
wherein the first member of a binding pair and the second member of a binding  
pair are independently selected from the group consisting of antigen, antibody,  
hormone, hormone receptor, polynucleotide, avidin, streptavidin, biotin, enzyme,  
enzyme substrate or inhibitor, lectins, specific carboxyhydrate, lipids, lipid binding  
10    proteins or membrane associated proteins, polynucleotides, polynucleotide  
binding proteins, receptor, transmitter, drug, target, protein, small molecule  
having a molecular weight of about 2,000 or less, polynucleic acid, DNA, and  
RNA.

15           7.     The surface plasmon resonance system according to claim 3,  
wherein at least one of the first member of a binding pair and the second  
member of a binding pair have a molecular weight of about 2,000 or less.

20           8.     The surface plasmon resonance system according to claim 1,  
wherein the second medium comprises a first member of a binding pair  
immobilized to the metal film in the assay region.

25           9.     The surface plasmon resonance system according to claim 1,  
wherein the metal film comprises at least one of gold, silver, aluminum, and  
molybdenum, and the first medium comprises at least one of a plastic,  
borosilicate glass, phosphosilicate glass, and silica glass.

10.    A method of monitoring a binding event, comprising:  
          illuminating a metal film through a first medium adjacent one side of  
the metal film, an assay medium comprising a first member of a binding pair

immobilized on a second side of the metal film;

contacting the first member of a binding pair with a second member  
of a binding pair;

collecting illumination reflected from the metal film; and

5 analyzing properties of the collected illumination; wherein at least  
one of illuminating the metal film and collecting illumination comprises using an  
optical integrated circuit.

11. The method according to claim 10, wherein illuminating the metal  
10 film comprises using light having a wavelength from about 390 nm to about 2,500  
nm.

12. The method according to claim 10, wherein at least about 225  
assay media each comprising a first member of a binding pair are immobilized on  
15 the second side of the metal film.

13. The method according to claim 10, wherein analyzing properties of  
the reflected light determines at least one of a) measuring an amount of the  
second member of a binding pair bound to the first member of a binding pair in  
20 the assay, b) whether or not binding occurs between the first member of a  
binding pair and the second member of a binding pair, and c) distinguishing  
between specific and non-specific binding between the first member of a binding  
pair and the second member of a binding pair.

14. The method according to claim 10, wherein the optical integrated  
25 circuit uses a 2 x N optical switch to directing light at the metal film.

15. The method according to claim 12, wherein the assay media each  
have a volume of about 0.5  $\mu$ L or less.

16. The method according to claim 10, wherein at least one of the first member of a binding pair and the second member of a binding pair have a molecular weight of about 2,000 or less.

5

17. The method according to claim 10, wherein the first member of a binding pair and the second member of a binding pair are independently selected from the group consisting of antigen, antibody, hormone, hormone receptor, polynucleotide, avidin, streptavidin, biotin, enzyme, enzyme substrate or inhibitor, lectins, specific carboxyhydrate, lipids, lipid binding proteins or membrane associated proteins, polynucleotides, polynucleotide binding proteins, receptor, transmitter, drug, target, protein, small molecule having a molecular weight of about 2,000 or less, polynucleic acid, DNA, and RNA.

10

18. A microwell array for a surface plasmon resonance system, comprising:

15

a silicon substrate comprising a plurality of wells to accommodate an interaction between a first member of a binding pair and a second member of a binding pair;

20

a metal layer coupled to the silicon substrate so as to be positioned at the bottom of the wells;

an insulation layer adjacent the metal layer;

an optical integrated circuit connected to the insulation layer, the optical integrated circuit comprising at least one arrayed waveguide grating spectrometer and at least one optical switch, the optical integrated circuit configured to illuminate each of the plurality of wells.

25

19. The microwell array according to claim 18, wherein the silicon substrate comprises at least about 400 wells.

20. The microwell array according to claim 18, wherein the plurality of wells independently comprise at least one selected from the group consisting of antigen, antibody, hormone, hormone receptor, polynucleotide, avidin, streptavidin, biotin, enzyme, enzyme substrate or inhibitor, lectins, specific carboxyhydrate, lipids, lipid binding proteins or membrane associated proteins, polynucleotides, polynucleotide binding proteins, receptor, transmitter, drug, target, protein, small molecule having a molecular weight of about 2,000 or less, polynucleic acid, DNA, and RNA.

21. The microwell array according to claim 18, wherein the metal film comprises at least one of gold, silver, aluminum, and molybdenum, and the first medium comprises at least one of a plastic, borosilicate glass, phosphosilicate glass, and silica glass.

22. The surface plasmon resonance system according to claim 1, wherein the optical integrated circuit comprises an integrated optical switch.

23. The method according to claim 10, wherein the optical integrated circuit comprises an integrated optical switch.